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Antiferromagnetism in water doped $\text{YBa}_2\text{Cu}_3\text{O}_{6+x}$ for $x \sim 0.5$

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Abstract

We report zero-field (ZF) μSR on a series of antiferromagnetic (AFM) samples of $\text{YBa}_2\text{Cu}_3\text{O}_{6.4}$ reacted with water. We find several oscillating signals whose amplitudes vary with water content and temperature, and for high water contents, a non-oscillating component. The Néel transition is 395(5)K, characteristic of the undoped CuO_2 bilayer. Initial comparison with ZF μSR and ZFNMR measurements in related AFMs ($\text{PrBa}_2\text{Cu}_3\text{O}_7$ and H doped $\text{YBa}_2\text{Cu}_3\text{O}_7$) suggests a similar muon site and an AF structure which is similar, but distinct from $\text{YBa}_2\text{Cu}_3\text{O}_6$. © 2000 Elsevier Science B.V. All rights reserved.

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Reaction of water with the well-known cuprate superconductor YBCO_{6+x} for $x < 1$ leads to intercalation of H_2O or related species into the partially occupied chain layer yielding an insulating antiferromagnet [1,2]. The structure of the product of this reaction is an analogue of the 1248 superconductor with Cu vacancies in the (double) chain layer [3]. When the reactant is well-ordered ortho-II $\text{YBCO}_{6.5}$ (see Ref. [4] for recent refined structural studies), the resulting AFM has a very well-defined Cu ZFNMR spectrum which differs significantly from pure YBCO_6 (the AFI phase with an ordered moment of $\sim 0.6\mu_B$ on the plane copper (Cu(2)

[5]). Detailed studies of this compound are desirable because they amount to characterization a potentially common impurity phase,¹ and more significantly, a difference in AF structure which is suggested by the ZFNMR could be used to better determine the parameters governing the AF ordered phase and, as a corollary, the remnant antiferromagnetism which is important in determining the properties of the metallic state (including T_c [6]).

In order to understand the origin of the differences observed in ZFNMR [1,2] between this AFM and YBCO_6 , we performed μSR studies on

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¹ The reaction takes place (albeit slowly) at room temperature with ambient water vapour, even in conventional dessicators or for epoxy packed samples.